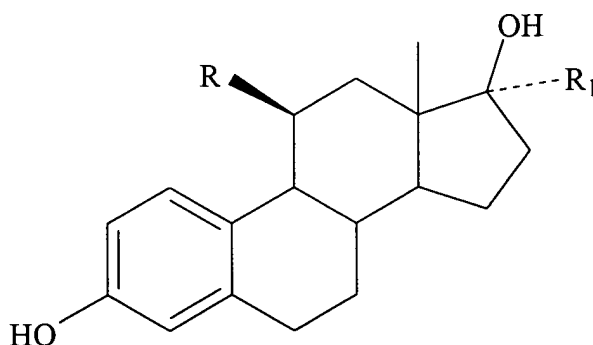


In the Claims:

Please amend the claims as follows.

1-38. Canceled.

39. (Currently amended) A method of treating the symptomology of menopause in a patient ~~at risk for developing~~ while reducing the risk that the patient develops an estrogen-sensitive cancer, the method comprising administering to said patient an effective amount of a compound selective estrogen receptor modulator (SERM) ~~according to~~ which has the chemical structure:



Where R is a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} Y R^1$  group, a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} R^2$  group, a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} R^3$  group, or a  $-(CH_2)_n X R^4$  group,

$R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each independently a  $C_1$ - $C_6$  linear, branch-chained or cyclo-alkyl group;

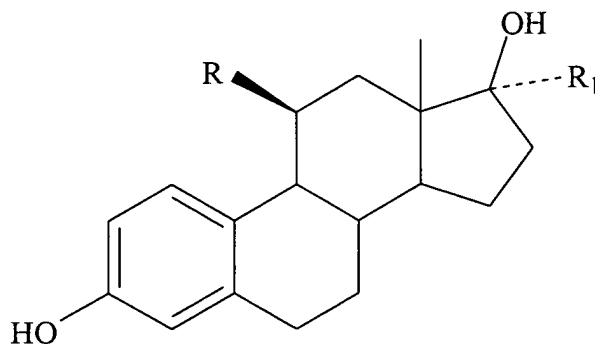
$R_1$  is H,  $CH_3$ , a vinyl group ( $-CH=CH_2$ ), or an ethynyl group ( $-C\equiv CH$ );

X is O or S and Y is O; and

n is from 1 to 3, wherein said symptomology is one or more of bone loss associated with osteoporosis, elevated cholesterol, elevated low-density lipoproteins (LDL) or cardiovascular disease associated with elevated cholesterol or elevated low-density lipoproteins (LDL).

40. (Previously presented) The method according to claim 39 wherein said menopausal symptomology is bone loss associated with osteoporosis.
41. (Currently amended)) The method according to claim 40 wherein ~~wherein~~ R is an ester or thioester group and R<sup>1</sup> and R<sup>2</sup> are each independently a C<sub>1</sub>-C<sub>5</sub> linear, branch-chained or cyclo- alkyl group.
42. (Previously presented) The method according to claim 39 wherein said compound is orally administered to said patient.
43. (Currently amended) The method according to claim 40 wherein said selective estrogen receptor modulator (SERM) is orally administered to said patient.
44. (Previously presented) The method according to claim 41 wherein X is O.
45. (Previously presented) The method according to claim 40 wherein X is O and R<sub>1</sub> is an ethynyl group.
46. (Previously presented) The method according to claim 40 wherein when R is an ester group and n is 1, and R<sup>1</sup> and R<sup>2</sup> have at least two carbon atoms.

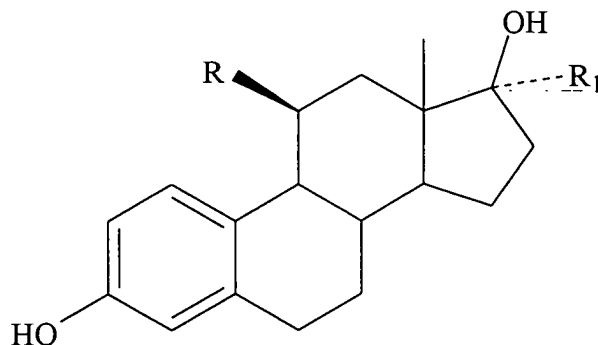
47. (Previously presented) The method according to claim 40 wherein when R is a keto, thioketo, ether or thioether group, n is 1, and R<sup>3</sup> and R<sup>4</sup> have at least three carbon atoms.
48. (Currently amended) A method of treating a patient suffering from an estrogen-sensitive cancer ~~in a patient~~, the method comprising administering to said patient an effective amount of a selective estrogen receptor modulator (SERM) ~~according to~~ which has the chemical structure:



Where R is a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} Y R^1$  group, a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} Y R^2$  group, a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} R^3$  group, or a  $-(CH_2)_n X R^4$  group,  
R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> are each independently a C<sub>1</sub>-C<sub>6</sub> linear, branch-chained or cyclo-alkyl group;  
R<sub>1</sub> is H, CH<sub>3</sub>, a vinyl group (-CH=CH<sub>2</sub>), or an ethynyl group (-C≡CH);  
X is O or S and Y is O; ~~and~~  
n is from 1 to 3; and  
wherein administration of the SERM does not exacerbate symptoms of the cancer and reduces the risk that the patient develops another estrogen-sensitive cancer.

49. (Previously presented) The method according to claim 48 wherein said estrogen-sensitive cancer is breast cancer.
50. (Currently amended) The method according to claim 49 wherein ~~wherein~~ R is an ester or thioester group and R<sup>1</sup> and R<sup>2</sup> are each independently a C<sub>1</sub>-C<sub>5</sub> linear, branch-chained or cyclo- alkyl group.
51. (Currently amended) The method according to claim 49 wherein said selective estrogen receptor modulator (SERM) is orally administered to said patient.
52. (Currently amended) The method according to claim 40 wherein said selective estrogen receptor modulator (SERM) is orally administered to said patient.
53. (Previously presented) The method according to claim 41 wherein X is O.
54. (Previously presented) The method according to claim 40 wherein X is O and R<sub>1</sub> is an ethynyl group.
55. (Previously presented) The method according to claim 40 wherein when R is an ester group and n is 1, and R<sup>1</sup> and R<sup>2</sup> have at least two carbon atoms.
56. (Previously presented) The method according to claim 40 wherein when R is a keto, thioketo, ether or thioether group, n is 1, and R<sup>3</sup> and R<sup>4</sup> have at least three carbon atoms.

57. (Currently amended) A method of reducing the likelihood of a recurrence of breast cancer in a patient ~~in need thereof~~ in remission from breast cancer, the method comprising administering to said patient an effective amount of a selective estrogen receptor modulator (SERM) according to which has the chemical structure:



Where R is a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} Y R^1$  group, a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} Y R^2$  group, a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} R^3$  group, or a  $-(CH_2)_n X R^4$  group,

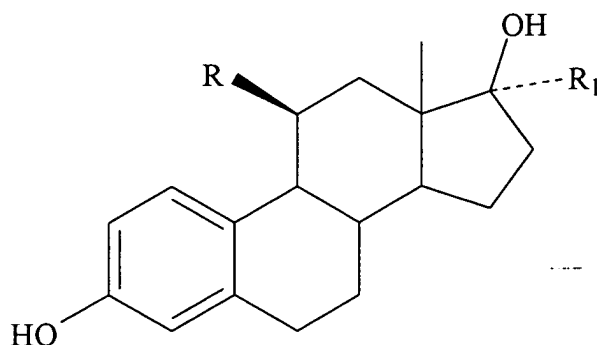
$R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each independently a  $C_1$ - $C_6$  linear, branch-chained or cyclo-alkyl group;

$R_1$  is H,  $CH_3$ , a vinyl group ( $-CH=CH_2$ ), or an ethynyl group ( $-C\equiv CH$ );

X is O or S and Y is O (~~preferably, X is O~~), and

n is from 1 to 3.

58. (Previously presented) The method according to claim 57 wherein R is an ester or thioester group and  $R^1$  and  $R^2$  are each independently a  $C_1$ - $C_5$  linear, branch-chained or cyclo- alkyl group.
59. (Previously presented) The method according to claim 57 wherein said compound is orally administered to said patient.
60. (Previously presented) The method according to claim 58 wherein said compound is orally administered to said patient.
61. (Previously presented) The method according to claim 57 wherein X is O.
62. (Previously presented) The method according to claim 57 wherein X is O and  $R_1$  is an ethynyl group.
63. (Previously presented) The method according to claim 57 wherein when R is an ester group and n is 1, and  $R^1$  and  $R^2$  have at least two carbon atoms.
64. (Previously presented) The method according to claim 57 wherein when R is a keto, thioketo, ether or thioether group, n is 1, and  $R^3$  and  $R^4$  have at least three carbon atoms.
65. (Currently amended) A method of treating the symptomology of menopause in a patient ~~with~~ suffering from an estrogen-sensitive cancer, the method comprising administering to said patient an effective amount of a selective estrogen receptor modulator (SERM) ~~according to which has~~ the chemical structure:



Where R is a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} Y R^1$  group, a  $-(CH_2)_n Y \overset{\overset{X}{\parallel}}{C} R^2$  group, a  $-(CH_2)_n \overset{\overset{X}{\parallel}}{C} R^3$  group, or a  $-(CH_2)_n X R^4$  group,

$R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are each independently a  $C_1$ - $C_6$  linear, branch-chained or cyclo-alkyl group;

$R_1$  is H,  $CH_3$ , a vinyl group ( $-CH=CH_2$ ), or an ethynyl group ( $-C\equiv CH$ );

X is O or S and Y is O; ~~and~~

n is from 1 to 3, wherein said symptomology is one or more of bone loss associated with osteoporosis, elevated cholesterol, or elevated low-density lipoproteins (LDL) or cardiovascular disease associated with elevated cholesterol or elevated low-density lipoproteins (LDL); and

wherein administration of the SERM does not exacerbate symptoms of the cancer and reduces the risk that the patient develops another estrogen-sensitive cancer.

66. (Previously presented) The method according to claim 65 wherein said menopausal symptomology is bone loss associated with osteoporosis.

67. (Previously presented) The method according to claim 65 wherein wherein R is an ester or thioester group and R<sup>1</sup> and R<sup>2</sup> are each independently a C<sub>1</sub>-C<sub>5</sub> linear, branch-chained or cyclo- alkyl group.
68. (Currently amended) The method according to claim 65 wherein said selective estrogen receptor modulator (SERM) is orally administered to said patient.
69. (Currently amended) The method according to claim 66 wherein said selective estrogen receptor modulator (SERM) is orally administered to said patient.
70. (Previously presented) The method according to claim 65 wherein X is O.
71. (Previously presented) The method according to claim 65 wherein X is O and R<sub>1</sub> is an ethynyl group.
72. (Previously presented) The method according to claim 65 wherein when R is an ester group and n is 1, and R<sup>1</sup> and R<sup>2</sup> have at least two carbon atoms.
73. (Previously presented) The method according to claim 65 wherein when R is a keto, thioketo, ether or thioether group, n is 1, and R<sup>3</sup> and R<sup>4</sup> have at least three carbon atoms.